Please substitute the following amended claims for corresponding claims previously presented. A copy of the amended claims showing current revisions is attached.

P

1. (Amended) A compound of formula I,

wherein

one of ${\sf R}^1$ and ${\sf R}^2$ represents a structural fragment of formula la

and the other represents R4;

Z represents O or N(R⁵);

 R^3 represents one or more optional substituents selected from OH, halo, cyano, nitro, C(O)OR⁶, C_{1.6} alkoxy or C_{1.6} alkyl (which two latter groups are

optionally substituted and/or terminated by one or more halo or hydroxy group)
or N(R⁷)R⁸;

 R^4 represents H, OH, halo, cyano, nitro, $C(O)OR^6$, $C_{1.6}$ alkoxy or $C_{1.6}$ alkyl (which two latter groups are optionally substituted and/or terminated by one or more halo or hydroxy group) or $N(R^7)R^8$;

Ar¹ represents phenyl, $C_{1\cdot3}$ alkylphenyl, $C_{1\cdot3}$ alkyldiphenyl, $C_{3\cdot7}$ cycloalkyl, $C_{1\cdot3}$ -alkyl- $C_{3\cdot7}$ -cycloalkyl, naphthyl, $C_{1\cdot3}$ alkylnaphthyl, thienyl, imidazolyl or isoxazolyl, all of which may be substituted by one or more substituent selected from OH, halo, cyano, nitro, $C(0)OR^6$, $C_{1\cdot6}$ alkoxy or $C_{1\cdot6}$ alkyl (which two latter groups are optionally substituted and/or terminated by one or more halo or hydroxy group) or $N(R^7)R^8$;

 R^5 represents H, $C_{1.6}$ alkyl, phenyl or $C_{1.3}$ alkylphenyl (which three latter groups are optionally substituted and/or terminated by one or more substituent selected from OH, halo, cyano, nitro, $C(O)OR^9$, $C(O)N(R^{10})R^{11}$, $P(O)(R^{12})R^{13}$, $P(O)(OR^{14})OR^{15}$, $S(O)_2(R^{16})R^{17}$, $S(O)_2N(R^{18})R^{19}$, $C_{1.6}$ alkoxy or $C_{1.6}$ alkyl (which two latter groups are optionally substituted and/or terminated by one or more halo or hydroxy group) or $N(R^{20})R^{21}$);

Y represents O, S, S(O), S(O)₂ or $N(R^{22})$;

 R^{10} and R^{11} independently represent H, OR^{23} , $C(O)R^{24}$, $OC(O)R^{25}$, $C(O)OR^{26}$, $C_{1\cdot4}$ alkyl, (which latter group is optionally substituted and/or terminated by one or more substituent selected from $C_{1\cdot4}$ alkyl, OR^{27} , $N(R^{28})R^{29}$, $C(O)OR^{30}$, $C(O)N(R^{31})R^{32}$, $P(O)(R^{33})R^{34}$, $P(O)(OR^{35})OR^{36}$ and $S(O)_2N(R^{37})R^{38}$), $-(CH_2CH_2O\cdot)_pR^{39}$ or, together with the nitrogen atom to which they are attached,

form a $C_{4.7}$ nitrogen-containing, aromatic or non-aromatic, ring which ring may contain a further heteroatom or group (as appropriate) selected from O, S and $N(R^{40})$ and may further be substituted by one or more substituent selected from $C(O)R^{41}$, $C(O)OR^{42}$ or $C(O)N(R^{43})R^{44}$;

 R^{28} , R^{29} , R^{30} , R^{31} , R^{32} and R^{40} independently represent H or $C_{1.6}$ alkyl, which latter group is optionally substituted and/or terminated by one or more substituent selected from $C(O)R^{45}$, $C(O)OR^{46}$ or $C(O)N(R^{47})R^{48}$; at each occurrence, R^6 , R^7 and R^8 independently represent H or $C_{1.4}$ alkyl; R^9 , R^{12} , R^{13} , R^{14} , R^{15} , R^{16} , R^{17} , R^{18} , R^{19} , R^{20} , R^{21} , R^{22} , R^{23} , R^{24} , R^{25} , R^{26} , R^{27} , R^{33} , R^{34} , R^{35} , R^{36} , R^{37} , R^{38} , R^{39} , R^{41} , R^{42} , R^{43} , R^{44} , R^{45} , R^{46} , R^{47} and R^{48} independently represent H or $C_{1.4}$ alkyl;

n represents 2;

p represents 1, 2, 3, 4, 5 or 6; and

B represents a structural fragment of formula lb, lc, ld or le

wherein



X¹ and X² independently represent a single bond or CH₂; or a pharmaceutically acceptable salt thereof.

 Ω^3

- 4. (Amended) A compound of formula I, as defined in claim 1, wherein R^2 represents a structural fragment of formula Ia and R^1 represents R^4 .
- 5. (Amended) A compound of formula I, as defined in claim 1, wherein Z represents O or $N(R^5)$, in which latter case R^5 represents $C_{1.6}$ alkyl terminated by $C(O)N(R^{10})R^{11}$.
- 6. (Amended) A compound of formula I, as defined in claim 1, wherein R³ is not present, or represents methyl, chloro or methoxy.

7. (Amended) A compound of formula I, as defined in claim 1, wherein Ar¹ represents substituted phenyl.

- 8. A compound of formula I, as defined in claim 1 wherein Y represents 0.
- 9. A compound of formula I, as defined in claim 1 wherein B represents a structural fragment of formula Ib.

23. (Amended) A pharmaceutical formulation including a compound as defined in claim 1, or a pharmaceutically acceptable salt thereof, in admixture with a pharmaceutically acceptable adjuvant, diluent or carrier.

- 24. (Amended) A compound as defined in claim 1, or a pharmaceutically acceptable salt thereof, for use as a pharmaceutical.
- 25. (Amended) A compound as defined in claim 1, or a pharmaceutically acceptable salt thereof, for use in the treatment of a condition where inhibition of thrombin is required.
- 26. (Amended) A compound as defined in claim 1, or a pharmaceutically acceptable salt thereof, for use in the treatment of thrombosis.

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- 27. (Amended) A compound of formula I as defined in claim 1, or a pharmaceutically acceptable salt thereof, for use as an anticoagulant.
- 28. (Amended) The use of a compound I as defined in claim 1, or a pharmaceutically acceptable salt thereof as active ingredient in the manufacture of a medicament for the treatment of a condition where inhibition of thrombin is required.

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- 30. (Amended) The use of a compound as defined in claim 1, or a pharmaceutically acceptable salt thereof, as active ingredient in the manufacture of an anticoagulant.
- 31. (Amended) A method of treatment of a condition where inhibition of thrombin is required which method comprises administration of a therapeutically effective amount of a compound as defined in claim 1, or a pharmaceutically acceptable salt thereof, to a person suffering from, or susceptible to, such a condition.